

ATTACHMENT A

World Trade Center Indoor Air Assessment: Selecting Contaminants of Potential Concern and Setting Health-Based Benchmarks

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Prepared by the Contaminants of Potential Concern (COPC) Committee
of the World Trade Center Indoor Air Taskforce Working Group

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Introduction

Since September 11, 2001, the outdoor environment around the World Trade Center (WTC) site and nearby areas has been extensively monitored by a group of federal, state, and local agencies¹. The agencies have taken samples of the air, dust, water, river sediments and drinking water and analyzed them for the presence of pollutants that might pose a health risk to response workers at the WTC site, office workers, and local residents. While some people may have experienced acute irritant and respiratory effects from the collapse of the towers and associated fires, extended monitoring of the ambient air at and beyond the perimeter of the WTC site over the past year indicates that contaminant concentrations pose a low risk of long-term health effects. During this same time period, limited investigation of the indoor environment in residential dwellings has occurred [NYCDOH, 2002; Chatfield & Kominsky, 2001]. As the cleanup of the WTC site is coming to an end, governmental health and environmental agencies are directing resources to evaluate the indoor environment for the presence of pollutants that might pose long-term health risks to local residents. Selecting Contaminants of Potential Concern (COPCs) and setting health-based benchmarks will assist the Pilot Cleaning Effectiveness Initiative² and inform the selection of contaminants in the Background Study³.

Selecting the Contaminants of Potential Concern (COPCs)

The assessment of the indoor environment began with a review of historical information on hazardous substances that have been associated with building fires and collapses [Wallace, 1990]. Many compounds, including combustion by-products such as dioxins and polycyclic aromatic hydrocarbons (PAHs) were identified for further investigation, along with building materials such as asbestos and fibrous glass. In addition, ambient air, indoor air, and indoor/outdoor bulk dust monitoring data were reviewed. Data sources included EPA's ambient air and bulk dust/debris monitoring program (www.epa.gov/wtc), OSHA's air/dust monitoring

¹ Agencies include the U.S. Environmental Protection Agency (EPA), Agency for Toxic and Disease Registry (ATSDR), New York State Department of Health (NYSDOH), New York City Department of Health (NYCDOH), Occupational Safety and Health Administration (OSHA), New York State Department of Environmental Conservation (NYSDEC) and New York City Department of Environmental Protection (NYCDEP).

²EPA is conducting a pilot program in an uncleaned/unoccupied building at 110 Liberty Street to determine the effectiveness of various cleaning methods for removing asbestos and other contaminants of potential concern from residential dwellings.

³Most if not all of the pollutants associated with the collapse of the World Trade Center were present in New York City's environment prior to September 11. To establish a baseline for the presence of these contaminants in affected residences, EPA will collect and analyze samples to look for some of these pollutants in apartments in parts of Manhattan that were not impacted. The Agency will use the data to determine pre-existing or "background" levels of these pollutants in interior spaces in New York City.

data and the NYCDOH/ATSDR indoor air pilot program [NYCDOH, 2002]. A concerted effort also was made to identify and review additional sources of WTC-related data from other governmental agencies (e.g., U.S. Geological Survey, NYC Board of Education), academic institutions, environmental organizations, and the private sector.

A semi-quantitative screening process was performed on the collected sampling data. Based on frequency-of-detection, concentration and inherent toxicity, contaminants that exceeded health-based screening levels for the ambient air (see Appendix B) were identified. Dioxin and PAHs were added to the COPC list by this process. In addition, building constituents with carcinogenic (asbestos) or irritant effects (fibrous glass, crystalline silica) that were consistently and significantly found in bulk debris and indoor dust samples were identified as COPCs [NYCDOH, 2002]. Finally, lead (Pb) was included based on a comparison of sampling data with existing regulatory standards. Collectively, the resulting group of contaminants are called “contaminants of potential concern” or COPCs in this report.

The list of COPCs is shown in the column at the left, below. Included for each COPC is a data summary that supports its inclusion on the list. For completeness, an additional list is included on the right containing substances that, for various reasons, have created concern, but on closer inspection, were either rarely detected in the ambient air and/or settled dust sampling around the WTC, or routinely found at concentrations below the health-based screening criteria presented in Appendix B.

<u>COPCs</u>	<u>Other Substances</u>
Lead	Benzene
PAHs	PCBs
Dioxin	Chromium
Asbestos	Cadmium
Fibrous Glass	Manganese
Crystalline Silica	Mercury
	Particulate Matter
	Refractory Ceramic Fibers
	Mold

Lead and Combustion By-Products (PAHs, Dioxin)

Lead. EPA’s ambient air monitoring data detected lead in concentrations that exceeded 1.5 ug/m³ on five occasions from September 18-27, 2001. It should be noted that the National Ambient Air Quality Standard (NAAQS) for lead is 1.5 ug/m³, but is based on a three-month average. Since the ambient monitoring program began, no three-month average of the ambient air data for lead exceeded the NAAQS of 1.5 ug/m³. However, lead was more commonly found at concentrations that exceeded 0.1 ug/m³, the default value employed in EPA’s Integrated Exposure Uptake Biokinetic Model for Lead in Children [EPA, 1994]. This value represents the upper-bound of average of ambient airborne lead concentrations in urban environments. Additional support for including lead as a COPC was provided by a report that characterized the

dust that settled over lower Manhattan after the WTC collapse [Lioy, 2002]. The lead concentration in settled dust ranged from 101 - 625 parts per million (ppm). By itself, this range does not represent extraordinarily high concentrations, but in combination with the mass of material released from the collapse of the towers, it represents a potentially significant increase in the mass of lead deposited in lower Manhattan. Therefore, lead is included as a COPC.

PAHs. Limited ambient air sampling was conducted for PAHs during the period of time (September through December, 2001) when combustion processes were occurring at the site. However, concentrations in bulk dust [EPA, 2002a and Lioy, 2002] exceeded the removal guidelines for soil in EPA's Hazard Evaluation Handbook [EPA, 1997] of 9 ppm based on benzo(a)pyrene equivalents. This value is risk-based, equating to an excess lifetime cancer risk of 1×10^{-4} over a 30-year exposure duration for residential settings. Therefore, PAHs are included as a COPC.

Dioxin. Ambient air samples exceeded EPA's screening criteria (see Appendix B) at various sampling stations during September and October, 2001. Data trend analysis for dioxin in ambient air [EPA, 2002a] showed that concentrations significantly exceeded the risk-based screening level of 0.001 - 0.005 ng/m³. This range was based on an excess lifetime cancer risk of 1×10^{-4} over a 30 year exposure duration for residential settings, using both the existing cancer Slope Factor [Integrated Risk Information System (IRIS), 2002] and proposed cancer Slope Factor in EPA's draft dioxin reassessment [EPA, 2001]. Therefore, dioxin (and related congeners) is included as a COPC.

Building Materials (Asbestos, Fibrous Glass, Crystalline Silica)

The evaluation of building materials for inclusion as COPCs required a different approach than that used for the combustion by-products. Those compounds (PAHs, dioxin) benefit from consensus toxicity criteria (i.e., IRIS-established cancer Slope Factors) that allow for a quantitative screening paradigm. With the exception of asbestos, the building materials evaluated (including fibrous glass, crystalline silica, mica, portlandite, calcite and gypsum) do not have IRIS inhalation toxicity criteria from which to perform quantitative risk assessments. Complicating this matter is the knowledge that many of these substances were deposited in a large cloud from the collapse of the towers. Thus, the settled dust may serve as a reservoir for re-suspension and eventual inhalation exposure. In addition, materials such as fibrous glass can also cause contact irritation when exposed directly to the skin.

The potential for exposure to building materials in the indoor environment was assessed by reviewing sampling data on the constituent composition of the indoor/outdoor bulk dust and indoor/ambient air [EPA, 2002c; USGS, 2001; NYCDOH, 2002; Chatfield & Kominsky, 2001].

Toxicity criteria for building materials (with the exception of asbestos) were obtained from occupational sources, such as the Occupational Safety and Health Administration's permissible exposure limits (OSHA PELs), and the American Conference of Governmental Industrial Hygienists threshold limit values (ACGIH TLVs). The evaluation of exposure potential coupled

with toxicity information identified three COPCs (asbestos, fibrous glass and crystalline silica).

Asbestos. In the days immediately following the disaster, EPA analyzed the settled bulk dust/debris for asbestos content. More than one third of the approximately 150 samples registered an asbestos concentration greater than 1% [EPA 2002c]. Since that time, EPA has taken the position that WTC-related dust should be considered potentially asbestos-contaminated and handled accordingly. Additionally, indoor settled dust samples collected from residential buildings in November and December 2001 indicated that 18% of the indoor locations sampled contained measurable levels of asbestos [NYCDOH, 2002]. Therefore, asbestos is included as a COPC.

Fibrous Glass. Analysis of WTC bulk dust/debris has consistently identified fibrous glass to be a major constituent of the material [Lioy 2002, USGS 2001]. In addition, an NYCDOH/ATSDR study [NYCDOH, 2002] found fibrous glass in the interior settled dust in 41% of the locations sampled at concentrations up to 35%. Air samples collected in areas with fibrous glass in settled dust indicate no fiber levels of immediate concern. Although fiber counts were found in four areas with slightly greater than background (0.004-0.006 f/cc), subsequent re-analysis indicated actual fibrous glass concentrations from these areas as 0.00004 to 0.00026 f/cc. Air samples from remaining areas showed a maximum 0.003 f/cc total fiber count by PCM. These wools may be skin, eye, and respiratory tract irritants. Although there are no standards to evaluate the settled dust content, the presence of fibrous glass in settled dust does indicate a potential for exposure. Therefore, fibrous glass is included as a COPC.

Crystalline Silica. Settled dust and air samples taken in indoor and outdoor areas of residential buildings in November and December of 2001 indicate the presence of alpha-quartz. Other forms of crystalline silica were not found. This is consistent with outdoor dust and debris samples collected by the USGS [USGS, 2001] and subjected to mineral analysis. Quartz was found in approximately 49% of the settled dust samples from indoor areas of residential buildings and all of the associated outdoor areas sampled. Levels of quartz ranged as high as an estimated 31.4% of the dust by weight in a residence. Since quartz is a common material in sand, finding this mineral in a city where there is a great deal of concrete is not unusual. However, quartz in dust from a comparison area unaffected by the WTC collapse ranged from non-detect only up to an estimated 2.2% in the residence [NYCDOH, 2002]. Seventeen residential areas and eleven common areas had quartz levels greater than the associated comparison area. Therefore, quartz was deemed to be elevated in some indoor areas of lower Manhattan relative to the comparison area. Additionally, quartz was found in 13% of the respirable fraction air samples taken in these areas, ranging from an estimated 4-19 ug/m³, demonstrating a potential for exposure. Although below occupational standards, this estimated concentration is above the effective NAAQS standard for the silica fraction of respirable particulate matter. Therefore crystalline silica, measured as alpha-quartz, is included as a COPC.

In addition to crystalline silica, calcite, portlandite and gypsum were the most abundant minerals detected in settled dust samples from residential areas in lower Manhattan following the WTC collapse. Mica was detected with much less frequency, generally at less than 0.1% of the dust.

Halite (salt) was also detected at trace levels. Calcite, portlandite, and gypsum are typical components of concrete and gypsum based wallboard products, which were present in the WTC buildings. While high concentrations of these minerals in airborne dust constitute a short-term health concern in the form of eye, nose and throat irritation, persisting adverse health effects would not be anticipated, unless these minerals remained suspended in high concentrations. Indoor and street-level outdoor air sampling done in November and December of 2001 show that the levels of these chemicals, over a time-weighted sample, were below levels associated with irritant effects (See Table 1). Although there are methodological difficulties in quantifying these materials by X-ray Diffraction (XRD) techniques, a semi-quantitative screening of the results showed that the air levels were orders of magnitude below occupational standards for irritant effects. However, since the presence of minerals in settled dust indicates there is potential for the materials to be resuspended, these minerals should be evaluated qualitatively. Therefore, although no screening levels are developed for these minerals as COPCs, the Pilot Cleaning Effectiveness Initiative mentioned in the Introduction will report qualitatively on the presence of gypsum, calcite and portlandite, while the quantitative analyses for crystalline silica (a COPC discussed above) are being performed by XRD (following NIOSH Method 7500).

Table 1. Index of NIOSH and OSHA exposure limits and estimated maximum values in Lower Manhattan.

Mineral	NIOSH REL (ug/m ³)	OSHA PEL (ug/m ³)	*Maximum Estimated Value (J) in Lower Manhattan (ug/m ³)
Gypsum	10,000 ug/m ³ (total) 5,000 (ug/m ³ resp)	15,000 ug/m ³ (total) 5,000 (resp)	14J (PM100) 15J (PM4)
Portlandite	5,000 ug/m ³	15,000 ug/m ³ (total) 5,000 ug/m ³ (resp)	95J (PM100) 84J (PM4)
Calcite	10,000 ug/m ³ (total) 5,000 ug/m ³ (resp)	15,000 ug/m ³ (total) 5,000 ug/m ³ (resp)	14J (PM100) 10J (PM 4)
NIOSH = National Institute of Occupational Safety and Health, Centers for Disease Control and Prevention REL = recommended exposure level/limit OSHA = Occupational Safety and Health Administration PEL = permissible exposure limit. resp = respirable * [NYCDOH, 2002]			

Other Substances that Were not Selected to be COPCs

The following summary is provided for substances that were investigated, but not included on the COPC list. A full discussion of the reasons is provided in Appendix A.

Chemical	Reason
Benzene	Benzene is very volatile and dissipates into ambient air quickly. Grab samples that recorded high (above the OSHA PEL of 1 ppm) concentrations on the debris pile consistently dropped to below detection limits (20 ppb) at the site perimeter.
PCBs	In over 500 ambient air samples taken around the WTC work zone, none exceeded EPA's screening level of concern for ambient air (see Appendix B).
Chromium	All measurements (over 300 samples) of chromium in ambient air around the WTC work zone have been below EPA's screening level of concern for the ambient air (see Appendix B). Measurements in settled dust have been below EPA Removal Action Guidelines [EPA, 1997].
Cadmium	Cadmium was found in only two out of 300 air samples at levels greater than EPA's screening level for ambient air (Appendix B). Measurements in settled dust have been below EPA Removal Action Guidelines [EPA, 1997].
Manganese	Manganese was found in only four of over 300 air samples at levels that were greater than EPA's screening level for ambient air (Appendix B). Measurements in settled dust have been below EPA Removal Action Guidelines [EPA, 1997].
Mercury	Concern regarding mercury has been raised by a report of slightly elevated blood mercury levels in four Port Authority Police Officers at the WTC site (69 were screened). A NYC Department of Design and Construction report [Rottner, 2002] concluded that air and bulk data have not shown elevated Hg levels. The report also noted that routine urine screening of NYC firefighters assigned to the site did not indicate any sign of Hg exposure. A U.S. Geological Survey leachability study [USGS 2001] of indoor and outdoor WTC settled dust samples showed that it is unlikely that harmful levels of mercury could ever be released to the air. In addition, indoor air sampling using a Lumex Analyzer at numerous residential dwellings close to Ground Zero determined mercury vapor concentrations to be well below EPA's Reference Concentration (RfC) of .3 ug/m ³ [Johnson, 2002].
Particulate Matter (PM)	Ambient monitoring in the WTC area has shown that, since late October, levels of PM 10 and PM 2.5 (two commonly measured inhalable PM sizes) have been below EPA's level of concern for the ambient air (see Appendix B), indicating that there is no continuing source of PM entering homes and offices. EPA's Environmental Data Trend Report World Trade Center, 9/11/01 - 4/24/02 [EPA, 2002a] concluded that PM 2.5 and PM 10 appear to have returned to levels of a steady background state.

Chemical	Reason
Refractory Ceramic Fibers (RCF)	Ceramic fiber-containing materials are not expected to be present in the WTC buildings, as they are mainly used in industrial high temperature applications. Limited sampling conducted by NYCDOH, 2002 did not indicate any ceramic fibers in four indoor air samples analyzed by scanning electron microscopy (SEM).
Mold	Mold was not directly associated with the WTC collapse. However, it may have resulted from the combination of water damage and inoperable HVAC systems. Guidelines for identifying and removing mold are provided through a NYCDOH website (see Appendix A).

Setting Benchmarks for the Contaminants of Potential Concern (COPCs)

A tiered approach was used to evaluate the health risks posed by contaminants that might be present in the indoor environment (air and settled dust). For each COPC, three levels were established:

- Tier I Level above which, after elimination of potential indoor sources (combustion by-products, stored chemicals, etc.), aggressive clean-up action should be taken expeditiously along with follow-up sampling to confirm attainment of Tier III level
- Tier II Range where diligent cleaning should continue, after elimination of potential indoor sources (combustion by-products, stored chemicals, etc.), with follow-up sampling to confirm attainment of Tier III level
- Tier III Level below which the risk is negligible or consistent with the New York City background level found in the aforementioned Background Study (see Introduction).

The following hierarchal approach was employed for developing benchmark values: 1) Use of relevant and appropriate environmental standards/regulations; 2) Calculation of risk-based benchmarks using EPA risk assessment guidance; and, 3) Adaptation of occupational standards with additional safety factors. Accordingly, a review of environmental standards/regulations was conducted for each of the six COPCs. As a result of this exercise Tier I screening levels for lead in indoor air and settled dust were set using EPA's National Ambient Air Quality Standard (NAAQS) and the U.S. Housing and Urban Development's (HUD's) standard for floors, respectively. The clearance criteria established in the Asbestos Hazard Emergency Response Act (AHERA, 1986) of 70 structures/mm² was utilized to evaluate asbestos samples from the ambient air monitoring effort. As detailed in Appendix B, the upper bound estimate for exposure duration to contaminants in the ambient air was one year. However, given the potential for

extended exposure in residential dwellings, AHERA was deemed less appropriate to this setting.

In cases where appropriate standards did not exist (e.g., asbestos), risk-based criteria were developed using established EPA risk assessment methods: for indoor air, methods described in EPA's "Risk Assessment Guidance for Superfund" [RAGS, 1989] were used; for settled dust, EPA Region III's "Wipe Sample Assessment" guidance was utilized with modifications (see Appendix D). The risk-based criteria reflect the most current toxicity criteria (Cancer Slope Factors and RfDs/RfCs) on EPA's Integrated Risk Information System (IRIS), which is a regularly updated (quarterly), online database that reports chemical toxicity reference values and information on human health effects that may result from exposure to chemicals in the environment.

For contaminants that lacked IRIS verified toxicity criteria, occupational standards were employed. Additional safety factors were added to account for higher exposure and greater sensitivity within the general population.

The clearance criteria paradigm described above is pathway and chemical specific. Accordingly, the benchmarks for each COPC do not account for multimedia and multiple chemical exposure.

Individual sampling results that exceed benchmark values should not be interpreted to represent the occurrence of an adverse health effect. Because benchmark levels assume continuous exposure for an extended duration, the average of the measured concentrations is more appropriate for evaluating risk than individual measurements. Consequently, isolated individual values above the benchmark level may not necessarily be indicative of a hazard.

Developing Risk-Based Criteria for Indoor Air

For carcinogenic compounds, the benchmarks were set so that a local resident's lifetime risk of developing cancer from exposure to WTC-related contaminants would not exceed a one-in-ten-thousand probability (1×10^{-4}) above the resident's background risk without this exposure. The amount of time that residents and office workers were exposed to WTC-related contaminants is unknown. To be conservative, the Tier I screening level was chosen to be protective of a resident who may have been exposed to WTC-related contaminants for one year. The Tier III clearance level was chosen to be protective of a resident who is exposed to WTC-related contaminants for 30 years, which is the upper-bound estimate for residency in one dwelling [EPA, 1989]. Cancer risk from less-than-lifetime inhalation exposure is given as:

$$\text{Risk} = \text{LAC} * \text{UR}$$

where LAC is the air concentration averaged over a lifetime, calculated as: $\text{AC} * [\text{EF} * \text{ED} / \text{LT}]$, where AC is the average air concentration during the period of exposure (g/m^3), EF is the exposure frequency (days/year), ED is the exposure duration (years), LT is lifetime (days), and UR is the unit risk factor, expressed in units of $1/\text{concentration}$.

The following table lists the input parameters and numerical values (along with a brief explanation) used in this procedure. (See Appendix D for equations and a more detailed discussion).

Table 2. Exposure parameters for calculating clearance criteria for air samples.

Input Parameter	Value	Explanation
Risk	1×10^{-4}	See Appendix C
AT-(Averaging Time - Carcinogens) of LT	25,550 days	See Appendix D
ED (Exposure Duration)	30 years	Upper-bound Estimate of Time in Residence [EPA, 1989]
EF (Exposure Frequency)	365 days/yr	Days in Residence [EPA, 1989]
IUR (Inhalation Unit Risk)	2.3 E -01 (asbestos) 2.86 E +5 (dioxin)* 7.3 E 0 (PAHs)*	IRIS, 2002 Dioxin Reassessment, EPA, 2001 IRIS, 2002

*IUR was calculated by route-to-route extrapolation of oral Slope Factor

The cancer risk level (1×10^{-4}) employed herein reflects the upper bound of the acceptable risk range (10^{-4} to 10^{-6}) established in EPA's Superfund regulation [Federal Register, 1990]. Practical Quantitation Limits (PQLs, the lowest level that can be reliably achieved within specified limits of precision and accuracy during routine laboratory operating procedures [EPA, 1992]) and anticipated background levels [ATSDR, 1995] dictated the selection of the risk level at 1×10^{-4} . A more detailed discussion of this subject can be found in Appendix C.

For non-carcinogenic compounds, airborne contaminant concentration is compared to the Reference Concentration (RfC). The RfC is an estimate of a chronic exposure concentration for the human population, including sensitive subpopulations, that is likely to be without an appreciable risk of deleterious effects during a lifetime [EPA, 1989]. That comparison (i.e., chronic exposure concentration (CEC) divided by the RfC) is called the Hazard Quotient (HQ).

$$HQ = CEC/RfC$$

where CEC is the daily averaged air concentration, calculated as: $AC * [EF*ED/AT-NC]$, where AC is the average air concentration during the period of exposure (g/m^3), EF is the exposure frequency (days), ED is exposure duration (years), AT-NC is the averaging time for non-carcinogens, and RfC is the reference concentration, expressed in units of 1/concentration.

According to EPA guidelines [EPA, 1989], if the HQ is greater than one, there may be concern for potential health effects. Therefore, for Tier III (clearance level), the benchmark screening

level was set at an HQ of 1. For Tier I, which equates to a sub-chronic exposure scenario, an HQ of 10 was used, in accordance with EPA's Hazard Evaluation Handbook [EPA, 1997]. An HQ of 10 accounts for the fact that chronic toxicity criteria (RfDs/RfCs) are being applied to sub-chronic exposure scenarios (i.e., exposure not expected to exceed 6 months to one year in duration). Accordingly, an HQ of 10 was used for non-carcinogens to reflect a similar (i.e., upper bound of 1 year) exposure duration. Note that contaminants (both non-carcinogens and carcinogens, alike) can exhibit acute effects from short-term, high-dose exposures. Because the Tier I benchmark levels are based on subchronic exposure (i.e., 1 year), acute effects from exposures that are below the benchmark levels would be unlikely. A review of EPA's draft Acute Exposure Guideline Levels [EPA, 2001a] and California EPA's (CAL-EPA) Acute Risk Levels demonstrates that the benchmark levels developed herein are categorically more stringent than the analogous US EPA and Cal-EPA acute levels.

Developing Risk-Based Criteria for Settled Dust

There is no established national guidance, and limited scientific literature, for assessing risk from interior surface wipe sampling (i.e., mass per unit area) data. The approach used in this report was adapted and modified from a methodology developed by the EPA Region III Superfund Program. It has been employed by the U.S. Army Corps of Engineers to develop risk-based clean-up goals for interior surfaces at the Claremont Polychemical Superfund site in Region II [Radian, 1999]. Additionally, the methodology is similar to an approach employed by the NYSDOH for evaluating a dioxin contaminated office building [NYSDOH, 1985] and one previously proposed by NJDEP for setting interior building surface clean-up goals [NJDEP, 1993].

A full discussion of the methodology employed for developing risk-based benchmarks based on contact with residues on indoor surfaces is presented in Appendix D. Briefly, this methodology provides an estimate of daily dose resulting from dermal absorption and ingestion associated with contacting contaminated surfaces and hand-to-mouth transfers. Dose is obtained by estimating the following: skin contact area and contact frequency with contaminated surfaces, transfer efficiency between surface and hand, transfer efficiency between hand and mouth, oral absorption of the contaminant and dermal absorption of residual contamination on skin. Consistent with the approach for indoor air, toxicity criteria were obtained from IRIS. The assessment procedure is described briefly below and in more detail in Appendix D.

For the purpose of developing risk-based clearance levels for interior surfaces, this approach is preferable to an approach that relies on measuring contaminant concentration on a mass per mass (ppm) basis (as might be done for an exterior soil assessment). After cleaning has occurred, it would be difficult to obtain the mass of dust necessary to perform a mass per mass measurement of contaminant concentration. In addition, literature on interior lead contamination demonstrates a strong correlation between lead load (mass per unit area) and blood lead concentration in children (Lanphear et. al., 1998).

The following table lists the input parameters and numerical values (along with a brief

explanation) used in this procedure. (See Appendix D for equations and a more detailed discussion.)

Table 3. Exposure parameters used to calculate clearance criteria for wipe samples.

Input Parameter	Value	Explanation
SA (Skin Surface Area)	400 cm ²	Palm Side of Hands
CF (Contact Frequency)	16/day	See Appendix D
FTSS (Fraction Transferred from Surface to Skin)	0.05	See Appendix D
EF (Exposure Frequency)	365 days/yr	Days in Residence [EPA, 1989]
ED (Exposure Duration)	30 years	Upper-bound Estimate of Time in Residence [EPA, 1989]
ABSo (Oral Absorption Fraction)	1	See Appendix D
FTSM (Fraction Transferred from Skin to Mouth)	0.1	See Appendix D
ABSd (Dermal Absorption Fraction)	0.03 (dioxin) 0.13 (PAHs)	Chemical Specific [EPA, 2001b]
BW (Body Weight)	70 kg	Average Adult [EPA, 1989]
AT-NC (Averaging Time - Non Carcinogen)	—	
AT-C (Averaging Time - Carcinogen)	25,550 days	See Appendix D
CSF (Cancer slope factor -dermal and oral)	1.0 E+6 (dioxin) 7.3 E 0 (PAHs)	Dioxin Reassessment: EPA, 2001 IRIS, 2002

In light of the limited data for many of the input parameters used to estimate dose from exposure to residues on surfaces, conservative exposure estimates were employed. Still, there may be a concern for highly exposed sub-populations. Infants and young children constitute such a group due to age-specific activities such as crawling and mouthing and increased surface-area to body-weight ratios. In cases where non-carcinogenic effects are driving clearance levels, the difference in dose on a mg/kg/day basis between an adult and a young child could be considerable. However, in this document the two compounds (dioxin, PAHs) for which the surface contact methodology has been used to develop clearance levels are carcinogens, so it is this effect that drives the clearance levels. As such, the Tier III clearance level is strongly influenced by overall exposure duration, which in this assessment has been set at 30 years.

Consequently, weighted over 30 years, the impact of a high-exposure period (i.e., early childhood) is diluted. It is acknowledged that Tier I (one year exposure duration) screening levels for carcinogens may underestimate cancer risk for children, but Tier I contaminant levels are not expected to persist, therefore the magnitude of any associated uncertainty would be small. If a contaminant was to be added to the COPC list that had its clearance level based on a non-carcinogenic endpoint, a separate wipe assessment analysis should be performed taking into account childhood exposure parameters. A more complete discussion of the uncertainties associated with the methodology for surface contact assessment is presented in Appendix D.

With the exception of asbestos, the extremely limited air sampling data for specific building constituents in indoor air dictated that an assessment approach had to be based on historical data relating the concentration of fibers in air (fibers per cubic centimeters or f/cc) to the load in settled dust (fibers per square centimeters or f/cm²) [Millette and Hays 1994]. Although the empirical relationships (called K factors) are not rigorous enough to rely on for a firm health decision, they are used in this report to define what might constitute elevated building material counts that would need to be addressed on a near-term basis. This approach may overestimate potential exposures but, as such, is protective.

Developing Benchmark Levels Based on Occupational Health Standards

For fibrous glass and silica, risk-based methods were not available for indoor air due to lack of established environmental toxicity criteria. Therefore, benchmark levels were set based on occupational health standards established by the American Conference of Governmental Industrial Hygienists [ACGIH] (called “threshold limit values” or TLVs) and the Occupational Safety and Health Administration [OSHA] (called “permissible exposure limits” or PELs). TLVs/PELs are established to protect employees who might have been exposed to substances during an eight-hour day, for five days a week. For Tier I levels, TLVs/PELs were divided by ten to be more protective of residents who might be exposed to substances twenty-four hours a day, seven days a week. In addition, the TLVs/PELs were established to protect a healthy worker. For Tier III levels, TLVs or PELs were divided by 100 to be protective of a more diverse population that includes sensitive individuals such as children, older adults, and the infirm.

The following tables show the benchmark levels developed for each contaminant of potential concern. Each table is accompanied by risk equations, toxicity criteria and a summary of any contaminant-specific assumptions that were made in developing the benchmark values.

Table 4. Lead

Tier	Action	Lead (settled dust)		Lead (indoor air)	
		Level	Basis	Level	Basis
I	Aggressive cleaning. If levels persist, take additional action to reduce exposure.	>40 ug/ft ²	1) HUD standard for floors 2) Residential Lead Hazard Standard (TSCA Section 403)	>1.5 ug/m ³	NAAQS (intended to keep 99.5% of children below 30 ug/dl)
II	Maintain recommended cleaning methods. Consider additional monitoring.	40 ug/ft ² to 25 ug/ft ² (or background)	25 ug/ft ² is HUD screening level for floors	1.5 ug/m ³ to 1 ug/m ³ (or background)	1 ug/m ³ is the calculated value using EPA's IEUBK Lead Model for Children*
III	No further action	<25 ug/ft ² (or background)	Level could also be set at that found in background, or unaffected, areas, so that no increase in risk due to lead would be expected.	1 ug/m ³ (or background)	Level could also be set at that found in background, or unaffected, areas, so that no increase in risk due to lead would be expected.

* EPA developed the Integrated Exposure Uptake Biokinetic (IEUBK) Lead Model [EPA, 1994] to evaluate multimedia lead exposure to children in residential settings. EPA established a goal of attaining a 95% probability that blood lead levels in children be less than 10 ug/dl [EPA 1994a]. Setting the indoor air lead concentration at 1 ug/m³ (and using background concentrations for lead in water, diet, soil and dust) the IEUBK Lead Model estimates that 96% of the blood lead probability distribution falls below 10 ug/dl. See Appendix E for model run data files and graph of blood lead probability distribution.

Table 5. Polycyclic Aromatic Hydrocarbons (PAHs)

Tier	Action	PAHs (settled dust)		PAHs (indoor air)	
		Level	Basis	Level	Basis
I	Aggressive cleaning. If levels persist, take additional action to reduce exposure.	>9 mg/m ²	9 mg/m ² represents a 1x10 ⁻⁴ risk estimate for a 1 year exposure	>6 ug/m ³	6 ug/m ³ represents a 1x10 ⁻⁴ risk estimate for a 1 year exposure
II	Maintain recommended cleaning methods. Consider additional monitoring	9 mg/m ² to 0.3 mg/m ² (or background)	0.3 mg/m ² represents a 1x10 ⁻⁴ risk estimate for a 30 year exposure	6 ug/m ³ to 0.2 ug/m ³ (or background)	0.2 ug/m ³ represents a 1x10 ⁻⁴ risk estimate for a 30 year exposure
III	No further action	<0.3 mg/m ² (or background)	Level could also be set at that found in background, or unaffected, areas, so that no increase in risk due to PAHs would be expected.	<0.2 ug/m ³ (or background)	Level could also be set at that found in background, or unaffected, areas, so that no increase in risk due to PAHs would be expected.

Toxicity criteria for benzo(a)pyrene:

Oral Slope Factor = 7.3 E 00 (mg/kg/day)⁻¹ (IRIS, 2002)

(Inhalation cancer risk was calculated by route-to- route extrapolation of oral Slope Factor)

The following carcinogenic PAHs are evaluated as benzo(a)pyrene-equivalents [EPA, 1993]:

<u>Compound</u>	<u>Relative Potency</u>
Benzo(a)pyrene	1
Benz(a)anthracene	0.1
Benzo(b)fluoranthene	0.1
Benzo(k)fluoranthene	0.01
Chrysene	0.001
Dibenzo(a,h)anthracene	1
Indeno(1,2,3-c,d)pyrene	0.1

Table 6. Dioxin

Tier	Action	Dioxin (settled dust)		Dioxin (indoor air)	
		Level	Basis	Level	Basis
I	Aggressive cleaning. If levels persist, take additional action to reduce exposure.	>120 ng/m ²	120 ng/m ² represents a 1x10 ⁻⁴ risk estimate for a 1 year exposure	>0.03 ng/m ³	0.03 ng/m ³ represents a 1x10 ⁻⁴ risk estimate for a 1 year exposure
II	Maintain recommended cleaning methods. Consider additional monitoring	120 ng/m ² to 4 ng/m ² (or background)	4 ng/m ² represents a 1x10 ⁻⁴ risk estimate for a 30 year exposure	0.03 ng/m ³ to 0.001 ng/m ³ (or background)	0.001 ng/m ³ represents a 1x10 ⁻⁴ risk estimate for a 30 year exposure
III	No further action	<4 ng/m ² (or background)	Level could also be set at that found in background, or unaffected, areas, so that no increase in risk due to dioxin would be expected.	<0.001 ng/m ³ (or background)	Level could also be set at that found in background, or unaffected, areas, so that no increase in risk due to dioxin would be expected.

Toxicity criteria for dioxin:

Oral Slope Factor = 1.0 E 06 (mg/kg/day)⁻¹ [EPA, 2001]

(Inhalation cancer risk was calculated by route-to- route extrapolation of oral Slope Factor)

The table above reflects the proposed cancer Slope Factor for 2,3,7,8 TCDD and the toxicity equivalence (TEQ) paradigm for carcinogenic dioxin/furan congeners present in EPA's draft dioxin reassessment [EPA, 2001]. An equivalent table reflecting the previous IRIS-verified cancer Slope Factor for 2,3,7,8 TCDD would be represented by multiplying all listed values by a factor of six.

The methodology for estimating dose from wipe sampling data (see Appendix D) is consistent with the approach NYSDOH used in assessing office buildings in Binghamton, with modifications made to input parameters to reflect residential rather than occupational exposure. In addition, the Tier III benchmark equates to a lifetime exposure level which is approximately one tenth of that which results from current dietary intake of dioxin (65 pg/d). It is thus

reasonable to assume that any lower level of exposure would result in an inconsequential reduction in the current risk levels experienced by the general U.S. population.

Information regarding NYC background levels also would be an important consideration in determining the appropriate longer-term value. This report proposes selecting the longer-term value on the basis of a 1×10^{-4} risk level or NYC background, as determined by the WTC-directed Background Study.

Table 7. Asbestos

Tier	Action	Asbestos (Settled Dust)		Asbestos (indoor air)	
		Level	Basis	Level	Basis
I	Aggressive cleaning. If levels persist, take additional action to reduce exposure. Evaluate conditions in the area, including representativeness of air samples, asbestos settled dust levels, and loading.	> 30,000 f/cm ²	Millette's K factor to estimate airborne levels for different activities (see narrative below).	>0.028 f/cc (PCME*)	0.028 f/cc represents a 1×10^{-4} risk estimate for a 1 year exposure
II	Maintain recommended cleaning methods. Consider additional monitoring. Evaluate conditions in the area including representativeness of air samples, fibrous glass settled dust levels and loading.	30,000 f/cm ² to background	---	0.028 f/cc 0.0009 f/cc (or background)	0.0009 f/cc represents a 1×10^{-4} risk estimate for a 30 year exposure
III	No further action required.	Background	Level is set at that found in background, or unaffected, areas, so that no increase in risk due to the asbestos fibers would be expected.	0.0009 f/cc (PCME*) (or background)	Level is set at that found in background, or unaffected, areas. In this way, no increase in risk due to the asbestos fibers would be expected.

* PCME - fibers greater than 5 micrometers long (aspect ratio >3:1). Analysis by TEM.

Toxicity criteria for asbestos:

Inhalation Unit Risk = $2.3 \text{ E-01 (f/cc)}^{-1}$ [IRIS, 2002]

Risk-based criteria were used to develop the Tiered benchmark levels for asbestos in air. Conservative assumptions of continuous exposure to a constant level of airborne fibers for either 1 year (Tier I) or 30 years (Tier III) were combined with the IRIS Slope Factor to establish benchmarks representative of a 1×10^{-4} estimate of excess lifetime cancer risk for each scenario. This approach makes several assumptions, chief among those is the quantification of asbestos fibers in air based on the PCM definition of a fiber (greater than 5 μm in length with an aspect ratio of 3:1 or greater) and the use of the IRIS Slope Factor which was designed to apply to fibers so defined. Although there is some concern regarding shorter fibers, the approach used here represents the current consensus by the US EPA for quantifying risk of airborne asbestos fibers. It should be noted there is ongoing debate regarding the nature of health effects which may be attributed to shorter asbestos fibers. Both EPA and ATSDR are currently pursuing meetings to discuss and further refine these issues. However for the purposes of this response, addressing PCM equivalent fibers is considered protective.

The Tier I level for asbestos in settled dust was based on K factors [Millette and Hays, 1994], which are empirical relationships between concentrations of asbestos fibers in settled dust and indoor air. Millette developed the K factors by studying matched air and settled dust samples taken from various homes, at varying levels of activity in the home. The Tier I level was based on the K factor for a worst case scenario of a high level of activity. Although K factors are not rigorous enough to rely upon for a firm health decision, they are used here only to define what might constitute elevated fiber counts, that would need to be addressed as a near-term concern. This approach may overestimate potential exposures, but as such, is protective.

It should be noted that the airborne Tier I level of 0.028 f/cc PCM equivalents roughly equates to the AHERA TEM standard of 70 f/mm² (0.022 f/cc) in total fiber counts. Based on empirical information from the WTC ambient air monitoring program [EPA, 2002c] that recorded less than 20% of total AHERA TEM fiber counts to be >5 μm in length, the AHERA TEM standard of 70 f/mm² (0.022 f/cc) and the alternate AHERA PCM standard of 0.01 f/cc both meet the risk-based criteria established for the Tier I screening level.

Table 8. Fibrous Glass

Tier	Action	Fibrous Glass (settled dust)		Fibrous Glass (indoor air)	
		Level	Basis	Level	Basis
I	Aggressive cleaning. If levels persist, take additional action to reduce exposure. Evaluate conditions in the area including representativeness of air samples, fibrous glass settled dust levels and loading.	> 100,000 f/cm ²	Millette's K factor to estimate airborne levels for different activities (see narrative below).	> 0.1 f/cc	ACGIH TLV of 1 f/cc (see narrative below).
II	Maintain recommended cleaning methods. Consider additional monitoring. Evaluate conditions in the area including representativeness of air samples, fibrous glass settled dust levels and loading.	100,000 f/cm ² to background	---	0.01 to 0.1 f/cc	---
III	No further action required	background	Level is set at that found in background, or unaffected, areas. In this way, no increase in risk due to the fibers would be expected.	< 0.01 f/cc	ACGIH TLV of 1 f/cc (see narrative below)

* Fibers with length > 5µm with an aspect ratio > 3:1, as defined in the ACGIH TLV.

Air concentrations measured by PCM, NIOSH 7400, confirmed by SEM.

The benchmarks for airborne fibrous glass were set based on the consideration that low level exposures to fibrous glass would not pose the potential for significant long-term health effects given the expected low biopersistence of these materials [ATSDR, 2002]. The current occupational exposure standards for fibrous glass (glass and mineral wools) is established to prevent the upper respiratory tract and skin irritant effects. In order to apply this standard to a

general population and accounting for a longer daily exposure than seen in the occupation setting, uncertainty factors were used to reduce the allowable airborne levels for the Tiered benchmarks. The Tier I benchmark is set at one tenth of the TLV and the Tier III level is set at 1 hundredth of the TLV. It is believed these levels will prevent conditions which may cause irritant health effects for most individual.

Although fibrous glass in settled dust is known to be a contact irritant, little data exist to provide a threshold relating to what fiber loading may result in skin irritation, or airborne levels of concern. However, as with asbestos it was deemed preferable to have a Tier I benchmark available so that there was some upper bound to settled dust loadings that may trigger action. Here again Millet's K factors were used to correlate dust loading levels to the air benchmarks. The Tier I level for fibrous glass in settled dust was based on Millette's K factors, which are empirical relationships between concentrations of asbestos fibers in settled dust and indoor air. Millette developed the K factors by studying matched air and settled dust samples taken from various homes, at varying levels of activity in the home. The Tier I level was based on the K factor for a worst case scenario of a high level of activity. Although K factors are based on activity data for asbestos, they are believed to be protective for fibrous glass, which is larger and less airborne. In addition, although K factors are not rigorous enough to rely upon for a firm health decision, they are used here only to define what might constitute elevated fiber counts, that would need to be addressed as a near-term concern. This approach may overestimate potential exposures, but as such, is protective.

The Tier I level for fibrous glass in indoor air was based on the American Conference of Governmental Industrial Hygienists' (ACGIH) threshold limit value (TLV) of 1 f/cc. That TLV is based on limiting irritant effects of fibrous glass on workers. The TLV was divided by a factor of ten to account for the different exposure durations between ACGIH's workers and the WTC area's residents.

The Tier III level for fibrous glass in indoor air was based on the ACGIH TLV of 1 f/cc, divided by a factor of 100 to account for greater exposure and the different sensitivities between ACGIH's healthy works and the WTC area's more varied population.

Table 9. Crystalline silica - respirable fraction of alpha-quartz

Tier	Action	Crystalline Silica (settled dust)		Crystalline Silica (indoor air)	
		Level	Basis	Level	Basis
I	Aggressive cleaning. If levels persist, take additional action to reduce exposure. Evaluate conditions in the area including representativeness of air samples, asbestos settled dust levels and loading.	---	---	10.0 µg/m ³ *	OSHA PEL of 100 µg/m ³ (see narrative below).
II	Maintain recommended cleaning methods. Consider additional monitoring. Evaluate conditions in the area including representativeness of air samples, fibrous glass settled dust levels and loading.	above background		10.0 µg/m ³ to 1 µg/m ³ (or background) *	
III	No further action required.	background	Level is set at that found in background, or unaffected, areas. In this way, no increase in risk due to the silica in settled dust would be expected.	1 µg/m ³ or background	OSHA PEL of 100 µg/m ³ (see narrative below).

* Silica measured in airborne respirable dust, by NIOSH 7500.

The benchmarks for silica in indoor air were derived from the Occupational Safety and Health Administration's (OSHA) permissible exposure limit (PEL) for dust containing 100% respirable silica in air of 100 µg/m³ (0.1mg/m³) as a time weighted average for an 8 hour exposure.

Silica exposure poses risk of both cancer and non-cancer respiratory health effects. Although no Reference Concentration (RfC) or IRIS Slope Factor exists, there are methodologies for

examining risks of non-occupational exposures, on which to draw upon for a risk based approach to set benchmark levels. The approach taken here is a dual approach, where in the face of uncertainties the quantification of the benchmarks was based on the current occupational standards. Broader risk-based discussion of other methodologies provides support for the chosen benchmark levels although not quantitatively employed here to derive the benchmarks.

The Tier I benchmark is set at one tenth of the OSHA PEL. A single year of exposure at this benchmark ($10\mu\text{g}/\text{m}^3$) would provide a cumulative silica loading to the lungs less than the $1\text{mg}/\text{m}^3$ work-year loading which is believed to be the departure point for adverse health effects. Additionally this level is also the effective National Ambient Air Quality Standard for silica, where PM_{10} at $50\mu\text{g}/\text{m}^3$ contains no more than 10% crystalline silica. A recent review by supports the finding that this level would be protective of long-term non-cancer health effects [EPA, 1996]. The Tier III benchmark is set at one hundredth of the OSHA PEL. A 30 year continuous exposure at this level would not result in a cumulative silica loading to the lungs greater than the $1\text{mg}/\text{m}^3$ year loading which is believed to be the departure point for adverse health effects.

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APPENDIX A

Hazardous Substances Not Included in Indoor Environment Sampling Program

Benzene

Benzene is a volatile organic chemical (VOC) that was found in emissions from the open flames and smoldering debris left after the collapse of the WTC. The benzene may have originated from the airplanes' residual jet fuel or from the continuous burning of plastics and other materials contained within the collapsed towers.

Since the fires and smoldering debris persisted long after the collapse of the WTC towers, there is a possibility that harmful levels of benzene vapor could have accumulated in nearby indoor environments. However, because benzene is very volatile and dissipates into ambient air quickly, the vapor plumes would have had to contain high concentrations of benzene, and the indoor area would have had to be relatively close to the fire source in order for indoor amounts of benzene to build up to harmful levels. In addition, unlike dioxin, benzene cannot become adsorbed by dust particles and transported indoors to settle as a long-term potential hazard. Once the source of the benzene is extinguished completely from the area, then benzene will dissipate and will not be a hazard.

Results of samples taken *within* the immediate WTC work zone from September 11 through mid-January 2002, show benzene levels of up to 180,000 parts per billion (ppb). However, tests during the same time period *outside* the work zone show levels of benzene that were below the established EPA screening level of 20 ppb for the ambient air (see Appendix B), indicating how quickly benzene dissipates in the air. A specific example, for the day of October 3, is shown below:

<u>From debris pile at WTC North Tower</u>	<u>From debris pile at WTC South Tower</u>	<u>From Liberty & Greenwich Streets</u>
28,000-39,000 ppb	4,300 ppb	less than 20 ppb

From January 26 through April 23, 2002 (the most recent test date available) levels of benzene in ambient air have remained below the 20 ppb screening level, except for one grab sample from the immediate work zone on February 9 and another sample from the same location on February 12. These tests indicate that the source of harmful ambient benzene levels has been extinguished, and therefore there is no source of benzene available to accumulate in any indoor environment. Benzene will not be monitored in any further indoor environment testing.

Polychlorinated Biphenyls (PCBs)

PCBs were historically used as coolants and lubricants in transformers, capacitors, and other electrical equipment. It is possible that, with the WTC collapse, PCBs could have been dispersed into the ambient air and then transported indoors. However, in well over 500 ambient air samples taken around the WTC site, no measurements were found above EPA's screening level of 730 ng/m³ in the ambient air (see Appendix B). This indicates that PCBs in ambient air are not considered to be at harmful levels, therefore, PCBs will not be monitored in indoor

environment testing.

Chromium

Chromium is a naturally occurring element commonly used in metal alloys and plumbing coatings in high rise buildings such as the WTC. Chromium and its compounds can be found in air as very fine dust particles that eventually settle over land, and can cause cancer and other health problems if inhaled at high concentrations. To date, in over 300 air samples, levels of chromium measured in ambient air around the WTC work zone have all been below EPA's screening level of 0.6 ug/m^3 in the ambient air (see Appendix B). Measurements in settled dust [EPA 2002c, Lioy 2002] have been below EPA Removal Action Guidelines [EPA, 1997]. Because of these low values, chromium is not being considered a hazard at the WTC site. Therefore, chromium will not be monitored in any indoor environment testing.

Cadmium and Manganese

Cadmium and manganese, both hazardous air pollutants, were detected in some ambient air samples taken at the WTC site. However, in over 300 samples, cadmium was found in only two samples at levels over EPA's screening level of 0.2 ug/m^3 in the ambient air (see Appendix B). In over 300 samples, manganese was found in only four samples at levels above EPA's screening value of 0.5 ug/m^3 in the ambient air (see Appendix B). Because of the low ambient concentrations of these two metals, they will not be monitored in any indoor environment testing.

Mercury

In late December 2001, it was reported that four New York City Port Authority police officers were tested with elevated levels of mercury in their blood. This raised concerns over the air quality at the WTC area. The officers were retested under medical surveillance to determine whether the source of mercury was the air over the WTC. The retesting found that the levels of mercury in the officers' blood and urine were below levels of concern established by NYCDOH and the American Conference of Governmental Industrial Hygienists. The cause of the first elevated mercury levels is not known, but may be from something that the officers had eaten the day before the blood tests were done, and not WTC air. In fact, Rottner [2002] has reported only one mercury air sample above detection limits to date in the WTC area.

In addition, data from a U.S. Geological Survey leachability study [USGS, 2001] performed on both indoor and outdoor WTC dust samples found mercury in leachate samples only at trace levels (up to 18 parts per trillion [ppt] for outdoor dust and 130 ppt in indoor dust). Because only trace levels of mercury appear to be present in indoor dust samples, the probability that harmful levels could ever be released to the air is very low. Therefore, mercury will not be monitored in any further indoor environment testing.

Particulate Matter

Particulate Matter (as fine dust and smoke) in the 10 micron and smaller size range is inhalable and may cause throat and lung irritation. Because of the energy released by the WTC disaster, dust containing particulate matter in this size range was generated. If this dust had not been able to settle, or if there had remained a continuous source of particulates being released into the ambient air, then there would have been a concern that hazardous dust could infiltrate nearby homes and offices. However, ambient monitoring for fine particulate matter in the WTC area has shown that, since late October, the levels of PM 10 (inhalable dust fraction 10 microns and smaller) and PM 2.5 (respirable fraction 2.5 microns and smaller) have been below the levels of concern:

PM10	150 ug/m3	NAAQS 24-hour average
PM2.5	65 ug/m3	NAAQS 24-hour average
	40 ug/m3	Air Quality Index, 24-hour average (moderate impact)

This indicates that there is no continuing concern of hazardous fine particulate matter entering homes and offices in the WTC area. Therefore, particulate matter will not be monitored in any further indoor environment testing.

Refractory Ceramic Fibers (RCF)

Ceramic fibers are reasonably anticipated to be human carcinogens based on sufficient evidence of carcinogenicity in experimental animals [IARC V.43, 1988]. When administered by inhalation, rats of both sexes showed a significant increase in the incidence of benign and malignant tumors of the lung. However, there are no data available to evaluate the carcinogenicity of ceramic fibers in humans [IARC V.43, 1988].

It is not expected that ceramic fiber-containing materials were present in large quantities in the WTC buildings, since those materials are used mainly in industrial high temperature applications.

Some air sampling has been performed for ceramic fibers in a recently completed study by ATSDR and NYCDOH [NYCDOH, 2002]. During the period from November 4 through December 11, 2001, environmental samples were collected in 30 buildings in lower Manhattan and four locations above 59th Street. Settled dust samples were taken both indoors and out, and analyzed for asbestos and other fibers by PLM. Although fibrous glass was reported in many settled dust samples, RCF was not reported as a fiber present by PLM. Air samples, also taken in these areas, were analyzed for fibers by PCM. Six of these study area air samples appeared to contain fiber levels in air higher than the levels found above 59th Street. Four of the samples from areas containing fibrous glass in settled dust were re-analyzed by SEM and were not found to contain any ceramic fibers above the detection limit (0.00004 f/cc).

Even though this is a limited study, it appears to confirm limited to no use of ceramic fibers in

the WTC buildings. Therefore, ceramic fibers will not be monitored in any further indoor environment testing.

Mold

Mold was not generated during the WTC collapse or associated fires. However, mold may be present because of actions taken in response to the collapse. Substantial quantities of water were used to extinguish the fires and to wash buildings affected by the collapse. Some of that water entered buildings, causing indoor mold contamination. Additionally, the loss of electric power after the collapse affected the ability of buildings' ventilation and plumbing systems to control indoor environments. This may have contributed to indoor mold contamination in some buildings.

The most common symptoms of exposure to mold are runny nose, eye irritation, cough, congestion, and aggravation of asthma. Although there is evidence documenting severe health effects of mold in humans, most of this evidence is derived from eating mold-contaminated foods, or occupational exposures in agricultural settings, where inhalation exposures were very high. There are no numeric standards to define "safe" levels of mold. In addition, it is very unlikely that the existence of mold in buildings around the WTC site can be distinguished from mold conditions existing prior to the attack. Therefore, the guidance established below is qualitative in nature and targeted towards detecting the presence of mold and removing it, rather than quantifying levels that pose health risks.

When inspecting a building, investigators should:

1. Look for evidence of visible mold or water damage throughout the apartment or office. Pay particular attention to moldy odors.
2. Ask the building management if water damage was noted as a result of the WTC attack or events thereafter. The investigator should be alert for situations where the loss of power to a building may have created a situation where either the ventilation or plumbing systems failed or leaked, creating water damage.
3. If mold contamination is encountered, it should be removed in accordance with New York City Department of Health mold removal guidelines:

<http://www.nyc.gov/health/html/doh/html/epi/moldrpt1.html>

APPENDIX B

World Trade Center Health Effects Screening Criteria for Ambient Air

Introduction

Extensive air quality monitoring data have been collected at and around the World Trade Center (WTC) site since 9/11/01. Table 1 (Screening Criteria) is intended to provide health protective screening values for data evaluation. Analysis has been performed on an extensive list of potentially WTC-related contaminants. Many of the chemicals screened have demonstrated a consistently low (i.e., below detection limits or trace amounts) trend. Consequently, the list of contaminants in Table 1 represents those chemicals that, because of their intrinsic toxicity and frequency/magnitude-of -detection, pose the greatest potential hazard from exposure. This selection process (i.e., a toxicity/concentration analysis), although qualitative, reflects the contaminant-of-concern identification process recommended in the Risk Assessment Guidance for Superfund. Table 1 may be expanded as additional data analysis becomes available. Two populations have been identified for assessment: response/demolition (i.e., WTC site) workers; and residents living in Lower Manhattan (e.g., Battery Park City, Tribeca and other residential locations close to Ground Zero). Included in the resident category are all other workers located in Lower Manhattan with the exception of WTC site workers.

Relevant Standards

The following paradigm has been employed to develop screening values. For each of the two identified receptor populations (i.e., site workers and residents), existing standards are utilized where appropriate. Occupational standards (i.e., OSHA PELs) are used for all site workers conducting response/demolition activities covered by OSHA. Monitoring data from demolition areas are compared to OSHA PELs. (For example, the OSHA PEL of 1 ppm for benzene is used to evaluate benzene air samples taken directly from within the plume on the debris pile.) Environmental standards (e.g., NAAQS, AHERA) are utilized to evaluate monitoring data from the site perimeter and beyond where residents or non-WTC site workers may be exposed. (For example, lead air monitoring data from perimeter stations outside of the immediate work zone are evaluated against the NAAQS of 1.5 ug/m³.)

Risk-Based Screening Criteria

In cases where appropriate standards do not exist, risk-based screening criteria have been developed for residential (including the non-WTC site workers) receptors. (In the absence of OSHA standards, it is beyond the scope of EPA's mission to develop "occupational" screening values.) The risk assessment paradigm detailed in EPA's "Hazard Evaluation Handbook: A Guide to Removal Actions" (HEH) was employed for this initiative (except where otherwise noted in the Table 1 footnotes). Screening levels reflect the most current toxicity criteria (Slope Factors and RfCs) on EPA's IRIS database.

For carcinogenic compounds excess lifetime cancer risk was set at E-04 (one-in-ten thousand). The residential exposure scenario in the HEH was modified for carcinogens from the default of 30 years (upper-bound estimate for residency in one dwelling) to 1 year (to reflect an upper

bound estimate for the length of time a resident may be potentially exposed to WTC-related contaminants). In cases where the screening value based on a noncancer endpoint is more stringent, screening values for both cancer and noncancer endpoints are presented. It is also noted that the default 30 year exposure duration (and the 1 year site-specific adjustment) reflects an apportionment between child (20% of total exposure duration) and adult (80 % of total exposure duration) receptors. Because children have comparatively greater (as a function of body weight) respiration rates than adults, the screening values presented in Table 1 are marginally more stringent than values that would otherwise be derived by direct application of IRIS verified Unit Risk values.

For noncarcinogenic compounds, the Hazard Quotient (chronic daily intake/RfC) was set at 10. A Hazard Quotient of 10 is employed in the HEH to account for the fact that chronic toxicity criteria (RfDs/RfCs) are being applied to sub-chronic exposure scenarios that are not expected to exceed 6 months - 1 year in duration. Accordingly, a Hazard Quotient of 10 was utilized for non-carcinogens in Table 1 to reflect a similar (i.e., upper bound of 1 year) exposure duration. It is noted that contaminants (both non-carcinogens and carcinogens, alike) can exhibit acute effects from short-term, high-dose exposures. Because the screening values in Table 1 are based on subchronic exposure (i.e., 1 year), acute effects from exposures that are below the screening levels would be unlikely. Additionally, a review of California EPA's (CAL-EPA) Acute Risk Levels demonstrates that the screening criteria in Table 1 are categorically more stringent than the Cal-EPA's analogous acute levels.

NOTE: Individual sampling results that exceed screening values should not be interpreted to represent the occurrence of an adverse health effect. Rather, such information indicates the need for careful monitoring and the assessment of longer-term data trends for evaluation against appropriate health criteria. That is, most of the screening levels have been developed to account for continuous one year average exposure durations. Because these screening levels assume continuous exposure for an extended duration, the average of the measured concentrations is more appropriate for evaluating risk than an individual measurement. Consequently, miscellaneous individual values above the screening level may not necessarily be indicative of potential for concern.

Table 1
World Trade Center Screening Criteria

Contaminant	Site Worker ⁽¹⁾	Resident ⁽²⁾
<u>Inorganics</u>		
Asbestos ⁽³⁾	.1 f/cc (PCM)	70 S/mm2 (TEM)
Cadmium	5 ug/m3	.2 ug/m3 ⁽⁹⁾ 3 ug/m3 ⁽⁵⁾
Chromium ⁽⁴⁾	100 ug/m3	.6 ug/m3 ⁽⁵⁾

Contaminant	Site Worker ⁽¹⁾	Resident ⁽²⁾
Lead	50 ug/m3	1.5 ug/m3 ⁽⁷⁾
Manganese	5 mg/m3	.5 ug/3 ⁽⁶⁾
Sulfur Dioxide	5 ppm	.14 ppm ⁽⁷⁾
<u>Particulates</u>		
Total	15,000 ug/m3	NA
Respirable	5,000 ug/m3	NA
PM _{2.5}	NA	40 ug/m3 ⁽⁸⁾ 65 ug/m3 ⁽⁷⁾
PM ₁₀	NA	150 ug/3 ^(7,8)
<u>Semivolatiles</u>		
Dioxin/Furans (TEQ)	NA	.162 ng/m3 ⁽⁵⁾
PCBs	1,000 ug/m3	.73 ug/m3 ⁽⁶⁾ 9 ug/m3 ⁽⁵⁾
PAHs ⁽¹⁶⁾	NA	6 ug/m3 ^(5, 17)
<u>Volatiles</u>		
Acetone	1,000 ppm	1.5 ppm ⁽⁶⁾
Benzaldehyde	NA	860 ppm
Benzene	1 ppm	.02 ppm ⁽⁹⁾ .21 ppm ⁽⁵⁾
Benzonitrile	NA	NA
1,3 Butadiene	1 ppm	.01 ppm ^(5, 15)
Chloromethane	100 ppm	.4 ppm ⁽⁶⁾ 2.6 ppm ⁽⁵⁾
1,4 Dioxane	100 ppm	.5 ppm ⁽⁵⁾
Ethanol	1,000 ppm	45 ppm ⁽¹⁰⁾
Ethylbenzene	100 ppm	2.5 ppm ⁽⁶⁾
Freon 22	1,000 ppm ⁽¹⁴⁾	140 ppm

Contaminant	Site Worker ⁽¹⁾	Resident ⁽²⁾
Propylene	LEL ⁽¹³⁾	simple asphyxiant
Styrene	100 ppm	2.3 ppm ⁽⁶⁾
alpha methylstyrene	100 ppm	.1 ppm ⁽⁶⁾
Tetrahydrofuran	200 ppm	.9 ppm ⁽⁵⁾
Toluene	200 ppm	1.1 ppm ⁽⁶⁾
Xylenes	100 ppm	1 ppm ⁽¹¹⁾
<u>Reactive Gases</u>		
Acetaldehyde	200 ppm	.05 ppm ⁽⁶⁾ 1.3 ppm ⁽⁵⁾
Formaldehyde	.75 ppm	.04 ppm ⁽¹²⁾ .35 ppm ⁽⁵⁾
Acrolein	.1 ppm	.0001 ppm ⁽⁶⁾

Units

f/cc = fibers (>5 um length) per cubic centimeter of air

S/mm2 = structures (>.5 um length) per square millimeter of filter paper

ppm = parts per million in air

ug/m3 = micrograms of contaminant per cubic meter of air

ng/m3 = nanograms of contaminant per cubic meter of air

NA - Not Applicable

Footnotes:

1. “Site Workers” refers to all workers involved in the response/demolition of the World Trade Center. Listed values are Occupational Safety and Health Administration (OSHA) Permissible Exposure Limits (PELs), Time Weighted Averages (TWA) unless otherwise noted.

2. “Residents” refers to people living in the vicinity of the World Trade Center as well as all other potentially exposed workers not involved in the response/demolition

3. Resident screening value is based on Asbestos Hazard Emergency Response Act (AHERA) methodology which uses transmission electron microscopy (TEM), and because of its basis in “background” (vs a risk basis) includes all asbestos fibers greater than 0.5 microns in length. Worker values are based on phase contrast microscopy (PCM, - which doesn’t distinguish asbestos from other fibers) or, for results above the PCM screening value, TEM to derive a PCM

equivalence that includes all asbestos fibers greater than 5 microns in length.

4. Screening values for chromium were based on the most toxic form (hexavalent)
5. EPA - Hazard Evaluation Handbook (HEH) (carcinogen) > 1 year of continuous exposure equating to an excess lifetime cancer risk of one-in ten thousand
6. EPA - HEH (noncarcinogen) > Hazard Quotient (HQ) = 10
7. National Ambient Air Quality Standard (NAAQS)
 - Lead is a 3 month average
 - PM_{2.5} is a 24 hour average
 - Sulfur Dioxide is a 24 hour average primary standard
8. Air Quality Index (AQI)
9. Non cancer effects based on CAL-EPA toxicity studies
10. American Conference of Governmental Industrial Hygienists (ACGIH) Threshold Limit Value (TLV)
11. Agency for Toxic Substances and Disease Registry (ATSDR) Inhalation minimum risk level (MRL) x 10
12. ATSDR acute MRL
13. Lower Explosive Limit (2 - 11 %)
14. National Institute of Occupational Safety and Health (NIOSH)
15. Proposed Reference Concentration (RfC) - HEH (noncancer) > Hazard Quotient (HQ) = 10
16. Based on Benzo(a)pyrene toxicity equivalency factor toxicity equivalency factor (TEF)
17. EPA National Center for Environmental Assessment (NCEA) provisional inhalation Slope Factor (3.1 E 00 mg/kg/day⁻¹)

APPENDIX C

Basis for Tier III screening level of 1 E-04

Defensible analytical methodology and sampling protocols are being chosen for future indoor sampling and analysis activities. The methods chosen are ones that have been published by reputable agencies and are in common practice among testing laboratories. In some cases, minor modifications may be made to the sampling and analytical protocols, but these will be modifications that are well established in the laboratory community.

All protocols chosen are designed to reach the lowest level of detection that is reasonable for the established methods. For Dioxin, Asbestos and PAHs, the sampling and analytical protocols are designed to reach detection limits that represent risk estimate levels of 1 E-04. To reach risk estimates of 1E-06, extraordinary modifications would have to be employed. These modifications would either have to be incorporated into the analytical protocols to increase the sensitivity of the required instrumentation, incorporated into the sampling protocols to achieve a larger sample, or a combination of both. For the Chemical of Potential Concern (COPC) list, the analytical protocols chosen are already incorporating the maximum sensitivity of the instrumentation. Therefore, the only legitimate mechanism to lower the overall limits of detection is to modify the sampling protocol. The two means of achieving this goal are to either run the sampling equipment (pumps) at a higher flow rate, or for longer periods of time. For the COPC list modifying flow rates would involve operating the equipment to achieve flow rates on the order of 500 to 1000 liters per minute. The only equipment available to operate at such flow rates are large units that can not be brought inside a residence. Rates this high also present problems with creating excessive negative pressure for indoor environments, plus flow rates this high have not been tested using the sampling protocols, and there is high likelihood of having analyte breakthrough on the collection filters. Therefore, this is not practical. The other option is to run the equipment for long periods of time. Again with the list of Chemicals of Potential Concern, sampling periods of up to 800 hours (33 days of continuous operation) would be needed to reach the E-06 risk detection levels.

For silica, the analytical and sampling protocols chosen will give detection levels in the neighborhood of 5 ug/m³. Instrumental sensitivity can not be set any higher to reach lower detection levels. Also, the sampling protocols involved for this analysis have been thoroughly validated by NIOSH. Any change in pump flow rate or sampling duration beyond what is documented in the method will produce results that have not been validated. Therefore, the sampling protocol should not be changed from that which is documented.

For fibrous glass the methodology is such that detection levels as low as 0.00001 f/cc can be achieved. This is well below required levels of detection for future indoor studies.

Another consideration in setting the target risk level involved the anticipated background level of contaminants such as asbestos, dioxin and PAHs in urban indoor environments. As previously mentioned, EPA is currently conducting a study to characterize background conditions for WTC COPCs in New York City residential dwellings. In advance of this study, a literature review was conducted to provide a general estimate of background concentrations for carcinogenic COPCs in urban indoor environments. It should be noted that the literature is limited in this regard. For

asbestos, ATSDR reports that “measured indoor air values range widely, depending on the amount, type, and condition (friability) of asbestos-containing materials used in the building” [ATSDR, 1995]. In its review ATSDR notes that the studies suffer from lack of common measurement reporting units. Study results have been reported as ng/m^3 , f/cc (TEM) and f/cc (PCM). Using unit conversion factors recommended by the National Research Council in 1984, ATSDR [1995] reports that the arithmetic mean concentrations of monitoring data from a variety of indoor locations ranged from .00003 - .006 f/cc (PCM). The proposed clearance level for WTC-impacted residential dwellings (.0009 PCM equivalents) is within this background range.

APPENDIX D

Assessing Exposures to Indoor Air and to Residues on Indoor Surfaces

Introduction

The purpose of this Appendix is to provide further details on how procedures were selected to estimate exposure to indoor air and to residues on indoor surfaces in residences impacted by the WTC attack.

Indoor Air

Deriving clearance criteria for air samples was completed by using methods described in EPA's "Risk Assessment Guidance for Superfund" [RAGS, 1989]. These methods were developed to assess the risk from contaminants at Superfund sites. The clearance criteria were calculated using the formula below:

$$\text{Clearance criteria} = (TR \times AT) / (ED \times EF \times IUR)$$

where:

TR = Target Risk

AT = Averaging Time - Carcinogens

ED = Exposure Duration

EF = Exposure Frequency

IUR = Inhalation Unit Risk

Target Risk

The target risk identified for these calculations was 1×10^{-4} . Appendix C explains the rationale for this value.

Averaging Time - Carcinogens

For non-carcinogens, AT is the exposure duration expressed in days. For carcinogens, exposure is averaged over a 70-year lifetime (the factor on which the cancer slope factors are based), and the AT is 70 years, in days (25,550).

Exposure Duration

A value of 30 years is assumed to match upper bound estimate of time in a residence (EPA, 1997b).

Exposure Frequency

A value of 365 days/year is used to represent a full time resident.

Inhalation Unit Risk

The upper-bound excess lifetime cancer risk estimated to result from continuous exposure to an agent at a concentration of 1 g/m^3 in air. The interpretation of unit risk would be as follows: if unit risk = $1.5 \times 10^{-6} \text{ } \mu\text{g/m}^3$, 1.5 excess tumors are expected to develop per 1,000,000 people if exposed daily for a lifetime to $1 \text{ } \mu\text{g}$ of the chemical in 1 cubic meter of air. The inhalation unit risk values used in the calculations are $2.3 \text{ E-}01$ for asbestos (IRIS, 2002), $2.86 \text{ E} +05$ for dioxin (Dioxin Reassessment, EPA, 2001), and $7.3 \text{ E}+0$ for PAHs (IRIS, 2002).

Residues on Indoor Surfaces

The most formal EPA guidance which addresses this issue is the “Standard Operating Procedures (SOPs) for Residential Exposure Assessment” originally published by the Office of Pesticides in 1997 and updated in 2001 (EPA, 1997a and EPA, 2001a). This guidance was designed for estimating exposures to pesticides. Pesticides are typically applied to indoor surfaces as liquid or sprayed formulations which would create surface residues which are likely to be somewhat different than the fine dust particles associated with the WTC attack. So while this guidance was considered, a number of other sources were also reviewed including the Superfund guidance on dermal contact (EPA, 1989), the procedures used to develop re-entry guidelines for the Binghamton State Office Building (Kim and Hawley, 1985), procedures used by NJDEP for setting interior building surface clean-up goals (NJDEP, 1993), the building clean-up procedures presented by Michaud et al (1994) and an approach developed by the EPA Region III Superfund program that has been employed by the U.S. Army Corps of Engineers to develop risk-based clean-up goals for interior surfaces at the Claremont Polychemical Superfund site in Region II (Radian, 1999). Elements from these various methods were combined to derive the procedure shown below. The procedure involves 3 steps: 1) estimating the amount of chemical deposited on the skin using skin area, contact frequency and transfer efficiencies, 2) estimating the dermal dose based on an absorption fraction and 3) estimating the ingestion dose based on the fraction of material on hands that is transferred to the mouth.

1. Estimating Amount of Chemical Deposited on Skin

$$Ds = C \times SA \times CF \times FTSS \times EF \times ED$$

Ds	=	Amount of chemical deposited on skin (mg)
C	=	Concentration of chemical on contaminated surface (mg/cm ²)
SA	=	Exposed skin surface area (cm ²)
CF	=	Contact frequency of skin against surface (1/day)
FTSS	=	Fraction transferred from surface to skin
EF	=	Exposure frequency (days/year)
ED	=	Exposure duration (years)

2. Estimating Dermal Dose

$$Dd = (Ds \times (1 - FTSM) \times ABSd) / (BW \times AT)$$

Dd	=	Dermal dose (mg/kg/day)	BW	=	Body weight (kilograms)
FTSM	=	Fraction transferred from skin to mouth	AT	=	Averaging time (days)
ABSd	=	Dermal absorption fraction			

3. Estimating Oral Dose

$$Do = (Ds \times FTSM \times ABSo) / (BW \times AT)$$

Do = Oral dose (mg/kg/day)	BW = Body weight (kilograms)
FTSM = Fraction transferred from skin to mouth	AT = Averaging time (days)
ABSo = Oral absorption fraction	

The surface concentration term (C) was assumed to remain constant over the entire 30 year exposure period. This is probably not true for surfaces in the residences impacted by the WTC attack. Surface loadings will decline as a result of volatilization, chemical degradation, surface cleaning and transfers to skin/clothing. While some redeposition will also occur, the net long term effect should be a gradual decline. In this sense, the method will overestimate exposures. This same issue has been recognized in similar assessments involving building clean-ups and pesticide exposures:

- Two scenarios were considered in developing the procedures used to develop re-entry guidelines for the Binghamton State Office Building (Kim and Hawley, 1985). One assumed a constant surface concentration and the other assumed a first order exponential decay.
- The OPP guidance (EPA, 1997a and EPA, 2001a) uses a “dissipation” factor to account for degradation and other loss mechanisms after pesticide application. Similarly, Durkin et al (1995) has proposed a time-dependent transfer coefficient method for lawn treatment pesticides.
- Michaud et al (1994) proposed a mass balance model which accounts for losses from surfaces associated with building clean-ups.

Thus, while methods have been proposed to deal with changes in surface residue strength over time, it is uncertain how well these apply to the situation in residences near WTC. Thus for purposes of this screening assessment, it was decided to assume constant levels.

4. Calculating Clearance Criteria

In order to calculate a clearance criteria for wipe samples, the formulas identified above need to be rearranged to derive a concentration based on the target cancer risk or non-cancer hazard. The clearance criteria for settled dust, using a wipe sample, for each COPC can be calculated using the formula listed below and the exposure parameters listed in Table 3 in the body of the document.

$$\text{Clearance criteria (mg/m}^2\text{)} = (TR \times BW \times AT) / [(Dd \times CSFd) + (Do \times CSFo)] \times EF \times ED$$

The other terms in this procedure and associated uncertainties are discussed below:

Exposed Skin Surface Area (SA)

The skin surface area of 400 cm² is a typical value for the palm side of adult hands (EPA, 1997b) which is the body part that is most likely to contact surfaces.

Contact Frequency (CF)

Michaud et al (1994) assumed 8 contacts per day, apparently based on professional judgment.

Low-end values of 1/day and "worst-case" rates of 24/day (3/hour) may be assumed as a matter of professional judgment; such exposure would obviously involve a very physical job with frequent contact with the walls and floor. The value recommended here is 16 contacts per day and is based on professional judgement. If available, site-specific activity data should take precedence.

Fraction Transferred from Surface to Skin (FTSS)

This will vary depending on type of surface, type of residual, hand condition, force of contact, etc. Rodes et al. 2001 conducted experiments on particle transfer to dry skin and measured transfers of 10% from carpets and 50% from hard surfaces. These transfer efficiencies were found to decline with repeated contacts. The OPP guidance recommends 5% of application rate for carpets and 10 % for hard surfaces (EPA, 1997a). USEPA has previously assumed transfer of 0.5 for PCBs (EPA, 1987) based on an Office of Toxic Substances (OTS) assessment. Michaud et al (1994) assumed 0.5 for PCBs and dioxins, but stated that 0.1 might be more realistic. In developing re-entry guidelines for the Binghamton State Office Building after a fire, a 100% transfer was assumed (Kim and Hawley, 1985). In a study of Malathion uptake from different surfaces, USEPA-EMSL found that FTSS of malathion from painted sheetrock to human hands was only 0.0003. (Mean transfer from vinyl flooring to hands was 0.0018, and from carpet to hands was 0.0152.) Malathion is a pesticide assumed to have lipophilicity more similar to PCBs than to volatiles or metals. However, the representativeness of such a number for PCBs and dioxins is unknown. PCBs are more lipophilic (have higher K_{ow}s) than malathion. A value of 5% is recommended here. Although this is on the low end of the literature values, when combined with the high contact frequency, it provides a fairly high total transfer to skin.

Exposure Frequency (EF)

A value of 365 days/year is used to represent a full time resident.

Exposure Duration (ED)

A value of 30 years is assumed to match upper bound estimate of time in a residence (EPA, 1997b).

Fraction Transferred from Skin to Mouth (FTSM)

Michaud et al (1994) assumed that all of the residues deposited on the fingertips would be transferred to the mouth, twice per day. A similar approach is used in the OPP guidelines. In the Binghamton re-entry guideline derivation, a range of factors were used: 0.05, 0.1, and 0.25 representing the fraction of residue on hand that is transferred to the mouth (Kim and Hawley, 1985). A similar value of 10% is recommended here.

Oral Absorption Fraction (ABSO)

For chemicals whose dose-response parameters are based on experiments in which the absorption fraction is similar to the one expected in the exposure scenario, there is no need to adjust the RfD or CSF.

Body Weight (BW)

A value of 70 kilograms is assumed which represents an average adult (EPA, 1997b).

Averaging Time (AT)

For non-carcinogens, AT is the exposure duration expressed in days. For carcinogens, exposure is averaged over a 70-year lifetime (the factor on which the cancer slope factors are based), and the AT is 70 years, in days (25,550).

Dermal Absorption Fraction (ABSd)

This parameter is chemical-specific. Dermal absorption fractions of 0.06 for PCBs and 0.03 for dioxins from soil were first proposed in USEPA, 1992 and more recently adopted in EPA 2001b. Michaud et al (1994) used 0.02 for dioxins and 0.03 for PCBs uptake from a sooty surface, based on the ranges of estimated ABSd values for soil. The Binghamton panel used a range of values for PCBs (0.01, 0.1, and 0.5) and dioxins (0.01 and 0.1) (Kim and Hawley, 1985).

Reported ranges for dermal uptake for PCBs in solvent vehicles are reported to range from 15 to 56%, with most of the values clustering around 20% (ATSDR, 1993). Reported ranges for 2,3,7,8-TCDD in solvent vehicles are reported to range from 1 to 40% (ATSDR, 1988).

Therefore, it seems that even if absorption from the wall material might be enhanced by residual solvent, the maximum possible absorption of 100% would be unrealistic even for worst-case exposure.

The values recommended here of 3% for dioxins and 13% for PAHs are based on EPA, 2001b.

Toxicity Values

There are two toxicity values that can be used to calculate screening values, a Reference Dose (RfD) for non-carcinogenic compounds and a Cancer Slope Factor (CSF) for carcinogenic compounds. The RfD is defined as an estimate (with uncertainty spanning perhaps an order of magnitude) of a daily oral exposure to the human population (including sensitive subgroups) that is likely to be without an appreciable risk of deleterious effects during a lifetime. It can be derived from a NOAEL, LOAEL, or benchmark dose, with uncertainty factors generally applied to reflect limitations of the data used. The CSF is defined as an upper bound, approximating a 95% confidence limit, on the increased cancer risk from a lifetime exposure to an agent. This estimate, usually expressed in units of proportion (of a population) affected per mg/kg/day, is generally reserved for use in the low-dose region of the dose-response relationship, that is, for exposures corresponding to risks less than 1 in 100. As the cancer endpoint was more sensitive, CSFs were used instead of RfDs. The CSFs used to calculate the clearance criteria are: 1.0 E+6 for dioxin (Dioxin Reassessment, EPA, 2001) and 7.3 E+0 for PAHs (IRIS, 2002).

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APPENDIX E **IEUBK Model Results for Lead in Air**

LEAD MODEL FOR WINDOWS Version 1.0 Build 251

Model Version: 1.0 Build 251

User Name:

Date:

Site Name:

Operable Unit:

Run Mode: Research

The time step used in this model run: 1 - Every 4 Hours (6 times a day).

***** Air *****

Indoor Air Pb Concentration: 100.000 percent of outdoor.

Other Air Parameters:

Age	Time Outdoors (hours)	Ventilation Rate (m ³ /day)	Lung Absorption (%)	Outdoor Air Pb Conc (ug Pb/m ³)
.5-1	0.000	2.000	32.000	1.000
1-2	0.000	3.000	32.000	1.000
2-3	0.000	5.000	32.000	1.000
3-4	0.000	5.000	32.000	1.000
4-5	0.000	5.000	32.000	1.000
5-6	0.000	7.000	32.000	1.000
6-7	0.000	7.000	32.000	1.000

***** Diet *****

Age	Diet Intake(ug/day)
.5-1	5.530
1-2	5.780
2-3	6.490
3-4	6.240
4-5	6.010
5-6	6.340

6-7 7.000

***** Drinking Water *****

Water Consumption:

Age Water (L/day)

.5-1 0.200

1-2 0.500

2-3 0.520

3-4 0.530

4-5 0.550

5-6 0.580

6-7 0.590

Drinking Water Concentration: 4.000 ug Pb/L

***** Soil & Dust *****

Multiple Source Analysis Used

Average multiple source concentration: 240.000 ug/g

Mass fraction of outdoor soil to indoor dust conversion factor: 0.700

Outdoor airborne lead to indoor household dust lead concentration: 100.000

Use alternate indoor dust Pb sources? No

Age Soil (ug Pb/g) House Dust (ug Pb/g)

.5-1 200.000 240.000

1-2 200.000 240.000

2-3 200.000 240.000

3-4 200.000 240.000

4-5 200.000 240.000

5-6 200.000 240.000

6-7 200.000 240.000

***** Alternate Intake *****

Age Alternate (ug Pb/day)

.5-1 0.000

1-2 0.000

2-3 0.000

3-4 0.000

4-5 0.000
 5-6 0.000
 6-7 0.000

***** Maternal Contribution: Infant Model *****

Maternal Blood Concentration: 2.500 ug Pb/dL

CALCULATED BLOOD LEAD AND LEAD UPTAKES:

Year	Air (ug/dL)	Diet (ug/day)	Alternate (ug/day)	Water (ug/day)

.5-1	0.640	2.521	0.000	0.365
1-2	0.960	2.608	0.000	0.902
2-3	1.600	2.963	0.000	0.950
3-4	1.600	2.886	0.000	0.981
4-5	1.600	2.842	0.000	1.040
5-6	2.240	3.022	0.000	1.106
6-7	2.240	3.349	0.000	1.129

Year	Soil+Dust (ug/day)	Total (ug/day)	Blood (ug/dL)

.5-1	5.162	8.688	4.7
1-2	8.113	12.583	5.2
2-3	8.210	13.723	5.0
3-4	8.318	13.785	4.8
4-5	6.298	11.780	4.1
5-6	5.714	12.081	3.7
6-7	5.417	12.135	3.5

